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PATENT

**DEVICE AND METHOD FOR CONNECTING HOLLOW ORGANS
AND/OR SEALING WALL DEFECTS IN HOLLOW ORGANS**

The application claims priority of provisional patent application
60/297,127 filed June 8, 2001.

BACKGROUND OF THE INVENTION

The present invention relates to a device and a method for connecting hollow organs and/or sealing wall defects in hollow organs. Such devices and methods are used in the entire field of surgery for connecting hollow organs such as blood vessels or the bowel. These are used particularly in the field of vascular surgery, in particular in the field of bypass surgery in the case of coronary ischaemia for producing proximal or distal anastomoses or also for sealing vascular wall defects. They are used in particular for producing end to end, side to side or end to side anastomoses.

Nowadays, approximately 120,000 bypass operations are performed per year in Germany. The most difficult discipline amongst those are anastomoses on the coronary arteries. It demands a great deal of skill and experience of the surgeon as the vessels here are very small (average arterial diameter 2 mm, vein diameter approximately 4 mm, other transplant diameters for example mammae 2 mm) and no leakage or suturing to the vascular rear wall must occur.

The manual procedure can be divided into the following operational steps:

1. thoracotomy, sternotomy (opening and cutting through of the sternum);
2. transplant extraction and preparation of the transplant, for example a vein being used as the transplant;
3. putting in of the anastomatic suture with the individual steps of making an incision, preparation of the artery and connection of the transplant opening with the incision opening in the artery; and
4. closure of, the thorax.

In this procedure the connection of the blood vessels by the anastomotic suture represents the most difficult task.

Because there is a danger in this manual connection of suturing to the rear wall of the blood vessel or of leakage. In addition, operations of this type must take place generally on a non-beating heart, which implies additional traumatisation for the patient due to the use of the heart-lung machine.

DE 43 04 353 A1 and also US 5 545 148 A disclose an endoscopic suturing appliance with a helical needle which has one or more windings. This needle is guided in a rotating manner through the edges of a vascular opening which are both held together by clamps and situated opposite each other. In this manner, a spiral suture is put in, the needle pulling a thread through this suture. Finally, the thread ends are then suitably fixed or tied. The spiral needle is thereby freely guided and actuated by two rollers which are located on a handle-like mounting.

SUMMARY OF THE INVENTION

It is the object of the present invention to make available a device and a method for connecting hollow organs and/or for sealing wall defects in hollow organs, by means of which the suturing process can be performed more quickly and more simply and in particular in a high-quality form which is qualitatively reproducible.

This object is achieved by the device and the method disclosed hereinafter. Advantageous developments of the device according to the invention and of the method according to the invention are given as follows:

The device characterized in that the carrier element has suction openings for drawing in a tissue or a hollow organ.

The device is characterized in that the carrier element has an annular configuration.

The device is characterized in that the carrier element extends along the external edge of the first surface.

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The device is characterized in that the guidetrack is disposed along the recess in such a manner that the spiral needle can be guided at least partially between two edges of the recess which are situated opposite each other.

The device is characterized in that the guidetrack is disposed in portions along two edges of the recess which are situated opposite each other in such a manner that the portions of the guidetrack which are disposed along the edges of the recess which are situated opposite each other form segments of a single spiral.

The device is characterized in that at least two guidetracks are disposed in the base mounting and, situated opposite each other, extend in and introduced into the artery to be sutured. The probe knife has a probe at its tip which is flexible. In the central part of the probe knife there is located a blade which is aligned by the probe and hence the incision is made in the vascular wall taking into account for example the vascular wall thickness, plaque etc. At the same time, the opening of the base mounting can be orientated by rotation along the probe axis.

The recess in the base mounting has guidetracks which make it possible to position and to hold the transplant-side adapter in a defined manner above this recess. Instead of guidetracks, guidepins or the like or even magnets are possible.

Around the opening in the base mounting there can be located two channels on the underside of the base mounting which respectively form a semicircle and which represent guidetracks for spiral needles. These channels cut the opening such that a wall remains only laterally. On the underside of the wall there are further recesses of the guide tracks which serve as specific guide geometries for the spiral needles. Likewise around the opening, disposed between the recesses, there are located borings near the edge of the recess (opening) through which borings the edges of the incision can be drawn in and held in a defined manner. Further borings for drawing in the base mounting on the tissue are located distributed on the underside of the base mounting or on the annular part of a two-part base mounting.

The base mounting and/or transplant adapter are divisible in the center so that, after completion of the suturing, they can be withdrawn laterally respectively by one of their halves.

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The transplant-side adapter also has guidetracks which ensure exact positioning relative to the base mounting. Said adapter has in its center a receiving means (boring) which can draw in the transplant on its external side by means of suction openings which are disposed radially on the internal side of the boring. There are located in addition recesses between the borings through which recesses the spiral needles are guided in a combined state with the base mounting. In this case, the channels of the guidetracks for the spiral needles in the base mounting and this recess in the transplant-side adapter form a common guidetrack for spiral needles.

In the case of the method according to the invention and the device according to the invention, a needle in one variant pulls a thread behind itself, the ends of which are tied to each other in a suitable manner after completion of the suturing. However, a spiral needle is also conceivable which remains in the suture as a thread replacement. The needle can then also be made of biodegradable materials.

With the device according to the invention and the method according to the invention, in particular leakages of the suture and also suturing to the rear wall of the coronary artery can be avoided. Occlusions of the artery can thus be avoided. In addition, the base mounting stabilizes the coronary artery by suction of the coronary artery and of the tissue surrounding it so that if necessary an operation can be performed also on a beating heart. The device according to the invention is suitable in addition for minimally invasive use, it being able to be introduced in this case into the body via small accesses without opening up the thorax. In addition, the artery- and transplant-side part of the instrument, i.e. the base mounting and the adapter can be introduced into the body by means of a trocar and be operated endoscopically. The device according to the invention is also suitable for manipulation by a robot as long as the adaptation of the tool is configured appropriately.

The conception of the device according to the invention achieves the object of applying anastomoses (connection of hollow organs). Its field of application is first of all bypass surgery on the heart in the case of coronary ischaemia, however also in addition on other organs in the case of stenoses, occlusions, contractions and thromboses in peripheral arteries and generally for sealing openings in hollow organs. With the device according to the invention both

proximal and distal anastomoses can be applied in coronary artery bypass surgery (CABG, coronary artery bypass operation). Furthermore, applying an anastomosis can be effected with the mammae as the transplant.

BRIEF DESCRIPTION OF THE DRAWINGS

A few examples of devices and methods according to the invention are described subsequently.

Fig. 1 shows a base mounting according to the invention with adapter;
 Figs. 2a-b show the base mounting according to the invention;
 Figs. 3a-b show the adapter according to the invention;
 Figs. 4a-c shows three variants for a drive of a spiral needle;
 Fig. 5 shows a probe knife;
 Figs. 6a-c show the preparation of a transplant;
 Fig. 7 shows an adapter with transplant;
 Fig. 8 shows a base mounting with artery;
 Fig. 9 shows a side view of a base mounting and of an adapter;
 Fig. 10 shows a further section through a base mounting and an adapter;
 Figs. 11a-b shows suturing of an anastomosis;
 Fig. 12 shows a base mounting and an adapter;
 Figs. 13a-b show a further base mounting;
 Fig. 14 shows a section through a base mounting;
 Fig. 15 shows a view from below of the base mounting;
 Fig. 16 shows a divided base mounting; and
 Fig. 17 shows a device for sealing a vascular wall defect.

DESCRIPTION OF THE PREFERRED EMBODIMENT

Figure 1 shows a base mounting 1 according to the invention with adapter 2. The base mounting 1 is placed on an artery 4 and fixed by suction holes. By means of appropriate openings in the underside of the base mounting 1, which are not illustrated here, the base mounting 1 can now be drawn in and fixed to the heart

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muscle above the artery 4. The base mounting need not necessarily however have a channel-like recess 3. The artery can also be drawn in only after the heart muscle and via a separate suction device.

By means of a small cut in the artery, the configuration of the artery can now be determined by means of a probe, as is illustrated for example in Figure 5. The probe is provided with a cutting mechanism in order to make an incision in the artery along the probe direction, the incision being delimited by the opening 18 (see Figure 2) in the base mounting 1.

In addition, the base mounting has two guidetracks 11, 12 which are introduced as a boring in the base mounting 1. These guidetracks serve for guiding the spiral needles 12, 13. The actuation of the spiral needles is effected within the base mounting via the access 14, for example a flexible shaft.

An adapter 2 is introduced into the opening 18 of the base mounting 1 and has for its part an opening 5. A piece of a vein 6 is introduced as a transplant into the opening 5.

Both the base mounting 1 and the adapter 2 have one connection piece 7, 8 respectively to which low pressure tubes can be connected. In this way, a low pressure can be applied to corresponding openings along the circumference of the recess 18 or of the opening 15 in order to draw in and to fix the edge of the incision in the artery or the edge along the open end of the vein 6.

Figure 2 shows the base mounting 1 of Figure 1 on the one hand viewed from below (Figure 2A) and also in plan view (Figure 2B). Here, as in the following, identical or corresponding elements are provided with identical or corresponding reference numbers.

It can be detected in Figure 2A that the edge of the recess 18 has openings 15 on the one hand which represent guidetracks for a spiral needle. This means that the guidetracks 15 themselves are portions (segments) of a single spiral configuration so that a spiral runs through all of these segments 15 of the guidetrack during a rotating forward movement. Furthermore, both borings of the guidetrack 11 are illustrated which extend out with the recess 18 within the base mounting 1 and receive the spiral.

In Figure 2B, two guidepins 17 can be detected furthermore on the surface of the base mounting to the left and the right of the recess 18, which guidepins engage in corresponding borings in a guide element 9 of the adapter 2 and thus ensure a safe and correct fit of the adapter 2 in the base mounting 1.

Figure 3 shows the adapter 2 in both partial pictures A and B, in side view (Figure 3A) and in diagonal view (Figure 3B). The adapter 2 has a diagonally extending boring 5 into which a transplant can be introduced. On its lower end, which terminates approximately with the underside of the base mounting 1 in the inserted state, there are likewise provided guidetracks 20 for a spiral needle. These guidetracks are disposed in such a manner that they form with the guidetracks 15 in the base mounting a common guide for a spiral needle. Furthermore, there are located in this region suction openings 21 via which the edge of a transplant located in the opening 5 is drawn in and fixed to the external wall of the boring 5 in the adapter 2.

Figure 4 shows various possibilities for the drive of a spiral needle within the base mounting 1. The spiral needle 12 is located already within the spiral needle guide 10 in Figures A to C.

In Figure 4A, there are located beside the boring 10 in total three rollers 22 which are disposed in such a manner that their axis of rotation is located parallel to the boring 10. The rollers 22 are in addition disposed beside the boring 10 in such a manner that they remain in frictional contact in one of the spiral needles guided by the boring 10. If the rollers 22 are now actuated, then the spiral 12 is rotated and moves thereby forwards in addition.

A view is shown in Figure 4B in which in total five rollers 22 are used.

In Figure 4C, an actuation principle is chosen in which a shaft 23 is in non-positive contact with the rear end of the spiral needle. The shaft 23 is now rotated and at the same time pushed forwards so that the spiral needle is moved forwards while rotating.

Figure 5 shows a probe knife 30 with a handle 31 and a probe 33. Above the probe 33 in the direction of the handle 31 there is situated a knife edge 32 in the probe knife.

The probe can be introduced into a vessel and be aligned therein. If the probe 30 is aligned then an incision into the vessel can take place from inside by cutting through the vascular wall with the knife 32. In the case of an incision of this type, which is made within the recess 18 of the base mounting 1 into a vessel 3, the length of the incision is delimited by the size of the recess 18 in the base mounting 1.

Figure 6 shows the preparation of a transplant 6. For this purpose a transplant 6, for example a vein, is introduced into the opening 5 of the adapter 2. This is illustrated in Figures 6A and 6B. Next the transplant can be fixed to the wall of the boring 5 by suction, as is illustrated for example in Figure 3. The projecting portion of the transplant which projects downwardly over the adapter 2, is then cut off by a knife 34 along the lower edge of the adapter 2. This projecting portion 6 is then disposed of.

The proximal end of the transplant can however be effected via a type of template or limit stop corresponding to the incision on the artery.

Figure 7 shows the transplant 6 which is prepared and fixed in the adapter 2.

Next the base mounting 1 is positioned as illustrated in Figure 8 at a suitable position above a desired artery on the heart muscle and there is drawn in to the heart muscle via openings in the lower side of the base mounting 1 (not illustrated here).

Now the course of the artery is determined by a small cut by means of the knife probe 30. Advantageously, the base mounting 1 can be mounted rotatably and then be aligned correspondingly to the probe axis. This position is then fixed and the incision is made in the artery 4 along the probe direction within the recess 18 of the base mounting 1. The opening 18 in the base mounting 1 thereby delimits the size or length of the incision. Next the artery is drawn in and fixed in this position.

Figure 9 now shows how the transplant-side adapter 2 is placed on the base mounting 1. In order to achieve the exact position, pins 17 of the base mounting 1 are thereby used which correspond to the borings in the guide 9 of the adapter 2.

Next the spiral needles 12 and 13 are introduced into their corresponding needle guidetracks 10 or 11.

Figure 12 shows a further base mounting with adapter 2. Next to the borings 10 and 11 for guiding the spiral needles, rollers 22 are disposed respectively now underneath openings in the base mounting, said rollers being actuated via shafts 24. These rollers correspond to those from Figure 4A and serve for actuation of the spiral needles 12 and 13.

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CONCLUSIONS

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Figure 17 shows a further embodiment for sealing an opening in a hollow organ wall. The opening 40, for example an incision into an artery, is thereby likewise pierced by means of a spiral needle, the spiral needle going through both edges of the incision 40. After the spiral has passed through the edges of the incision 40, a thread is pulled through the thus pierced openings in the vascular wall by the spiral and then is tied.

In this case, a base mounting 1 occurs once again which has a guide 11 for a spiral needle 13. The spiral needle is thereby guided once again by the guide 11 in the region of an opening 18 partially within the base mounting and partially out with the base mounting in the recess 18. The channel 11, 15 thereby cuts the opening 18 such that there remains only a wall running laterally. On the underside of the wall, hollows can also be disposed which serve as guide geometries for the spiral needle. Here too, around the opening 18, for example between the recesses, borings can be provided close to the edge of the recess 18 through which the edges of the wall opening of the hollow organ can be drawn in and be held in a defined manner.

FIG. 17